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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/597,591	08/16/2006	Pann-Ghill Suh	20010-21USA	3560
35736 JHK LAW	I EX			
P.O. BOX 1078			MOHAMED, ABDEL A	
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			1654	
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			10/28/2008	PAPER

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

	Application No.	Applicant(s)		
	10/597,591	SUH ET AL.		
Office Action Summary	Examiner	Art Unit		
	ABDEL A. MOHAMED	1654		
The MAILING DATE of this communication app Period for Reply	pears on the cover sheet with the o	correspondence address		
A SHORTENED STATUTORY PERIOD FOR REPLY WHICHEVER IS LONGER, FROM THE MAILING DOWN THE MAILING DOWN THE MAILING DOWN THE MAILING DOWN THE MAILING THE MAILING THE METERS AND THE MAILING THE METERS AND TH	ATE OF THIS COMMUNICATION 36(a). In no event, however, may a reply be tinuity will apply and will expire SIX (6) MONTHS from the cause the application to become ABANDONE	N. mely filed the mailing date of this communication. ED (35 U.S.C. § 133).		
Status				
1) Responsive to communication(s) filed on 16 A 2a) This action is FINAL . 2b) This 3) Since this application is in condition for alloward closed in accordance with the practice under E	action is non-final.			
Disposition of Claims				
4) ☐ Claim(s) 1-20 is/are pending in the application 4a) Of the above claim(s) is/are withdray 5) ☐ Claim(s) is/are allowed. 6) ☐ Claim(s) is/are rejected. 7) ☐ Claim(s) is/are objected to. 8) ☐ Claim(s) 1-20 are subject to restriction and/or expressions.	wn from consideration.			
Application Papers				
9) The specification is objected to by the Examine 10) The drawing(s) filed on is/are: a) acc Applicant may not request that any objection to the Replacement drawing sheet(s) including the correct 11) The oath or declaration is objected to by the Ex	epted or b) objected to by the drawing(s) be held in abeyance. Se ion is required if the drawing(s) is ob	e 37 CFR 1.85(a). jected to. See 37 CFR 1.121(d).		
Priority under 35 U.S.C. § 119				
 12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). a) All b) Some * c) None of: 1. Certified copies of the priority documents have been received. 2. Certified copies of the priority documents have been received in Application No. 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)). * See the attached detailed Office action for a list of the certified copies not received. 				
Attachment(s) 1) Notice of References Cited (PTO-892) 2) Notice of Draftsperson's Patent Drawing Review (PTO-948) 3) Information Disclosure Statement(s) (PTO/SB/08) Paper No(s)/Mail Date	4) Interview Summary Paper No(s)/Mail D 5) Notice of Informal F 6) Other:	ate		

ELECTION/RESTRICTION

Restriction is required under 35 U.S.C. 121 and 372.

This application contains the following inventions or groups of inventions, which are not so linked as to form a single general inventive concept under PCT Rule 13.1.

In accordance with 37 CFR 1.499, applicant is required, in reply to this action, to elect a single invention to which the claims must be restricted.

Group I, claim(s) 1-13 and 20, drawn to a polypeptide comprising a W-rich peptide and a conservative variant or functional fragment thereof or a W-rich mimic thereof, a method of preventing inflammation in a subject comprising the steps of providing an inflammation preventing effective amount of the polypeptide according to claim 1 or a W-rich peptide mimic thereof to the subject in need thereof and a pharmaceutical composition comprising the polypeptide according to claim 1 or W-rich mimic thereof.

Group II, claims(s) 14, drawn to a method of treating arthritis in a subject comprising the steps of providing an inflammation preventing effective amount of the polypeptide according to claim 1 or a W-rich peptide mimic thereof to the subject in need thereof.

Group III, claim(s) 15, drawn to a method of treating an auto-immune disease in a subject comprising the steps of providing a therapeutically effective amount of the polypeptide according to claim 1 or a W-rich peptide mimic thereof to the subject in need thereof.

Group IV, claim(s) 16, drawn to method of preventing binding of Aβ42 to human neutrophils comprising contacting the neutrophil with the polypeptide according to claim 1 or a W-rich peptide mimic thereof.

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Group V, claim(s) 17, drawn to a method of treating Alzheimer's disease comprising administering a therapeutically effective amount of the polypeptide according to claim 1 or a W-rich peptide mimic thereof to the subject in need thereof.

Group VI, claim(s) 18 and 19, drawn to a method of identifying a FPR class receptor antagonist wherein the FPR class receptor is a FPRL1 comprising the steps of providing a cell having a FPR class receptor; contacting the cell with a candidate antagonist compound; and identifying the candidate antagonist compound as an antagonist compound if the candidate binds to a FPR class receptor and inhibits its activity.

The inventions listed as Groups I-VI do not relate to a single general inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons: the methods of Groups I-VI do not correspond to the same technical features and are not connected in design, operation or effect because they differ in method steps, parameters and reagents used, although, the five groups use the same compounds as recited above, and as such, the method of Group I is directed to a method of preventing inflammation in a subject comprising the steps of providing an inflammation preventing effective amount of the polypeptide according to claim 1 or a W-rich peptide mimic thereof to the

subject in need thereof. Group II is directed to a method of treating arthritis in a subject comprising the steps of providing an inflammation preventing effective amount of the polypeptide according to claim 1 or a W-rich peptide mimic thereof to the subject in need thereof. Group III is directed to a method of treating an auto-immune disease in a subject comprising the steps of providing a therapeutically effective amount of the polypeptide according to claim 1 or a W-rich peptide mimic thereof to the subject in need thereof. Group IV is directed to a method of preventing binding of Aβ42 to human neutrophils comprising contacting the neutrophil with the polypeptide according to claim 1 or a W-rich peptide mimic thereof. Group IV is directed to a method of treating Alzheimer's disease comprising administering a therapeutically effective amount of the polypeptide according to claim 1 or a W-rich peptide mimic thereof to the subject in need thereof.

With respect to Group VI, this group differs from Groups I-V in using different compounds for different purposes. Group VI is directed to an assay method of identifying a FPR class receptor antagonist while Groups I-V are directed to various methods of treatment and prevention of different diseases and/or conditions using basically the same compounds for different purposes. Thus, Groups I-V are methods of treatment or prevention while Group VI is an assay method. Therefore, the methods of Groups I-VI as recited above do not correspond to the same technical features and are not connected in design, operations or effects because they differ in method steps, parameters and reagents used and functions, and as such, the methods as grouped are independent and distinct, each from the other because they represent different technical

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features and different inventive endeavors. Thus, the Groups require different patent and literature search and as such Groups I-VI do not share the same technical features, the inventions do not relate to the same inventive concept.

SEQUENCE ELECTION REQUIREMENT

The various sequences disclosed in Groups I-V of claims 6-11 encompass peptides having different structures which are patentably distinct and/or independent, one from the other, and capable of independent use. Further, there is no sequence linking each with other, it is only consensus.

Therefore, the sequences are patentably distinct because they are unrelated sequences and each unrelated sequence is considered a separate and distinct product. For an elected invention drawn to either amino acid or polypeptide sequences, the Applicant must elect a **single** peptide sequence (See MPEP 803.04). Due to the increasing large size of sequence databases which must be searched and the increasing numbers of applications requiring sequence searches, it creates an undue burden on the Office to search more than a single sequence (product) per application. For these reasons, the requirement of 37 CFR 1.141 *et seq.* is no longer waived and Applicant is required to elect a **single** sequence for examination. Applicant is reminded that this is a **restriction requirement**, not an election of species as contended by Applicant. Thus, Applicant is required to elect a single disclosed sequence and/or provide a single subsequence within a disclosed sequence wherein the subsequence

for the elected is searched because the inventions do not relate to a single inventive concept.

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

Applicant is advised that the reply to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed (37 CFR 1.143).

CONCLUSION AND FUTURE CORRESPONDANCE

Claims 1-20 are subject to restriction and/or election requirement.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to ABDEL A. MOHAMED whose telephone number is (571)272-0955. The examiner can normally be reached on First Friday off.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Cecilia Tsang can be reached on (571) 272-0562. The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300.

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Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

Mohamed/A. A. M./ Examiner, Art Unit 1654

/JON P WEBER/ Supervisory Patent Examiner, Art Unit 1657